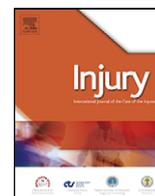




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2 Cadaveric assessment of a new guidewire insertion device for volar percutaneous
3 fixation of nondisplaced scaphoid fracture4 Marc Soubeyrand^{a,*}, Julien Even^a, Cesar Mansour^a, Olivier Gagey^a, Veronique Molina^a, David Biau^b5 ^a Hopital Universitaire de Bicetre, AP-HP, Bicetre F-94270, Univ Paris-Sud, Department of Orthopaedic Surgery, 78 rue du General Leclerc, 94270 Le Kremlin-Bicetre, France6 ^b Hopital Universitaire de Cochin, AP-HP, Department of Orthopaedic Surgery, 27 rue Fbg Saint Jacques, 75014, Paris, France

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ABSTRACT

Purpose: Volar percutaneous screw fixation (PSF) of acute nondisplaced scaphoid waist fractures allows early mobilisation of the wrist and a faster return to work than prolonged cast immobilisation. Usually, placement of the wire which guides the definitive cannulated screw is performed with the hand held. Nevertheless, correct placement of this wire is technically difficult. We designed a guidewire insertion device (GID) to facilitate this placement.

Methods: We compared the hand held technique with the technique using the GID in a cadaveric study. The hand held technique was performed on 16 scaphoids and the GID was used in 16 other scaphoids. The four participating surgeons were divided into two groups: two experienced surgeons and two inexperienced surgeons.

Results: The GID significantly decreased procedure duration ($P < 0.001$), number of attempts to place the wire ($P < 0.001$), and number of image-intensifier shots ($P < 0.001$). With both techniques, experienced surgeons were significantly faster ($P = 0.0083$) and required significantly fewer attempts ($P = 0.043$) than inexperienced surgeons. Using the GID, the procedure duration ($P = 0.0039$) and the number of image-intensifier shots ($P < 0.001$) decreased more with inexperienced surgeons than with experienced surgeons. As for the number of attempts, there was no statistical difference between the two groups ($P = 0.32$).

Conclusions: The GID decreased the time and radiation exposure needed to achieve correct volar percutaneous wire placement in the scaphoid, compared to the conventional hand held technique. Easier wire placement may lead surgeons to use PSF instead of prolonged cast immobilisation for treating nondisplaced scaphoid fractures.

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Introduction

9 The scaphoid is the most frequently fractured carpal bone and
10 the majority of such fractures occur at the waist.^{15,28} The annual
11 incidence of scaphoid fractures was estimated at about 43/100,000
12 overall¹⁸ and was markedly higher in men (38/100,000) than in
13 women (8/100,000).²⁰ Scaphoid fractures are managed by
14 physicians working in several specialties, including emergency
15 medicine, plastic surgery, and orthopaedic surgeons.^{3,9,23,26}
16 Scaphoid fractures have a major socioeconomic impact, as they
17 heal slowly and result in nonunion in about 12% cases,¹¹ leading to
18 prolonged work disability.^{6,31,32}

19 Acute nondisplaced scaphoid waist fractures are currently
20 treated with either cast immobilisation or percutaneous screw
21 fixation (PSF). PSF allows early mobilisation of both the wrist and
22 the elbow and a faster return to work, with comparable healing
23 times and rates but a lower overall cost, compared to cast
24 immobilisation.^{2,4,10,15,19,24,31} The first step of PSF is the insertion
25 of a guidewire, which is then used to guide the definitive
26 cannulated screw. Correct placement of this guidewire is crucial
27 to the success of the procedure but remains difficult to achieve, for
28 several reasons: the three-dimensional shape of the scaphoid is
29 very complex and it is a very small bone.^{5,7,8} Therefore, using two-
30 dimensional images for guidewire positioning offers limited
31 accuracy. However, considerable accuracy is required, since the
32 bone is small and incorrect guidewire placement increases the risk
33 of nonunion.¹ Although both anteroposterior and lateral radio-
34 graphs are used to position the guidewire, the surgeon can see only
35 one view at a time on the image intensifier. Adjusting guidewire
36 position in one plane without modifying its position in the other
37 plane is extremely difficult. Repeated adjustments are required to

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38 achieve correct placement in both planes, which increases both the
39 duration of the procedure and the radiation dose received by the
40 patient and surgeon. Methods for facilitating screw placement
41 have been developed. Herbert et al. designed a jig for screw
42 insertion during open surgery.¹⁶ A cadaver study suggested that
43 fluoroscopic navigation may help to achieve correct guidewire
44 placement.²¹

45 The first author of this manuscript designed an original guide
46 insertion device (GID) that allows adjustment of guidewire
47 position on one view (anteroposterior or lateral) without modify-
48 ing the position on the other view. The aim of this cadaver study
49 was to assess the impact of using GID on the easiness, duration and
50 success of completing the procedure.

51 **Materials and methods**

52 We used 32 scaphoids (16 left and 16 right) from 16 fresh frozen
53 cadavers. Each wrist was examined visually and fluoroscopically
54 for any evidence of trauma or surgery. Wrists showing such
55 evidence were to be excluded from the study.

56 Four surgeons with different levels of expertise in hand surgery
57 participated in the study: two were orthopaedic surgery residents
58 who had no experience with scaphoid PSF and two were senior
59 hand surgeons who had considerable experience with scaphoid
60 PSF. Each surgeon performed the procedure on eight scaphoids,
61 using the GID for two left and two right scaphoids and the hand
62 held technique for two left and two right scaphoids. Scaphoid
63 allocation to the guidewire device or hand held technique was
64 randomised and stratified on surgeon experience and side.

65 The GID was made of aluminum. Manufacturing plans were
66 created using 3DSMax® software (Autodesk, San Rafael, CA). The
67 device was composed of a horizontal base carrying two parallel
68 vertical blades separated by a 5-mm space (Fig. 1). A cylindrical
69 component 5 mm in diameter moved freely between the two
70 blades. The cylinder was cannulated to allow the passage of

71 guidewires 1 mm in diameter. This cylinder was called “reducer”
72 because it allowed to reduce the diameter of the wire’s pathway
73 from 5 mm (the space between the two blades) to 1 mm (the
74 diameter of the guidewire).

75 For both the hand held technique and the technique using the
76 GID, the upper limb was placed on an armboard. A Siremobil®
77 (Siemens, Munich, Germany) image intensifier set on zoom mode
78 was used. A 5-mm skin incision was made over the trapezium, and
79 the soft tissues were dissected down to the bone. A wire 1 mm in
80 diameter was put on a drill and the tip of the wire was inserted into
81 the tubercle of the scaphoid. The entry point was that described by
82 Menapace et al.²⁵ The wire was pushed into the scaphoid under
83 image-intensifier guidance, using anteroposterior and lateral
84 views, the goal being placing the guidewire in the centre of the
85 proximal pole without exiting the scaphoid. During the procedure,
86 the wire position was assessed on the image-intensifier views by
87 both experienced surgeons and one inexperienced surgeon. If at
88 least one of the experienced surgeons determined that the
89 guidewire position was incorrect, the guidewire was removed
90 and a new attempt at correct placement was made. The procedure
91 was stopped when the two experienced surgeons judged that
92 guidewire position was correct (Fig. 2). For each procedure, the
93 following data were recorded on a form: experience of the surgeon
94 (inexperienced/experienced), side of the wrist, duration of the
95 procedure, number of wire-placement attempts, and number of
96 image-intensifier shots.

97 **Hand held technique**

98 The C-arm of the image intensifier was positioned vertically.
99 The wrist was placed under the image-intensifier beam and could
100 be moved freely in pronosupination and flexion-extension to
101 obtain anteroposterior, oblique and lateral views of the scaphoid.
102 The guidewire was guided by hand and pushed with the drill into
103 the scaphoid.

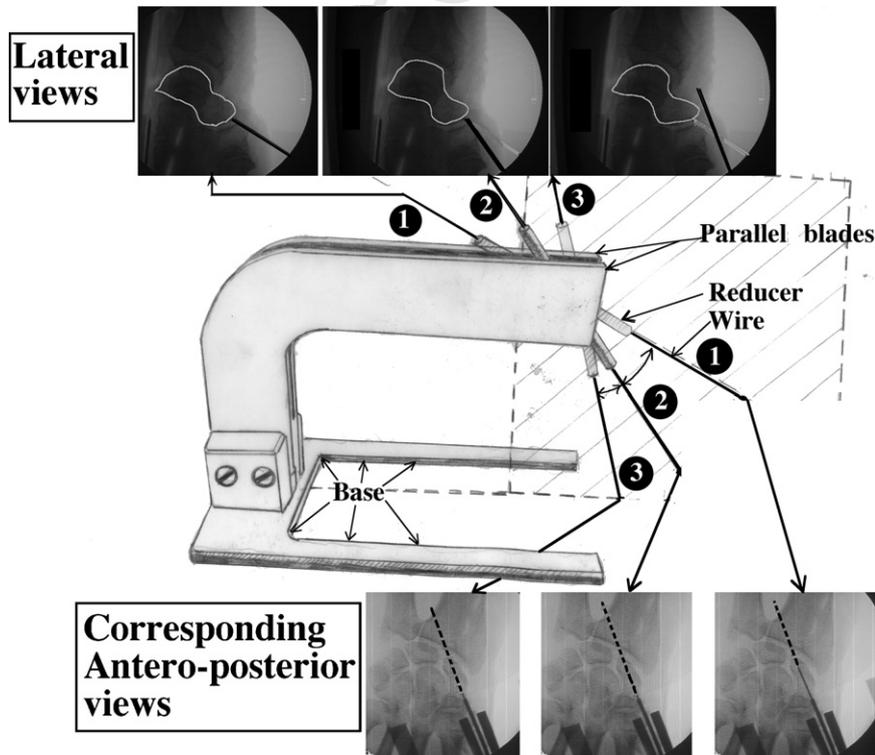


Fig. 1. Description of the GID. The wire is inside of the cylindrical reducer which is freely mobilisable between both the blades of the guidewire device. While the wire is mobilised between the blades, its direction on the lateral views changes while it remains the same on the anteroposterior views.

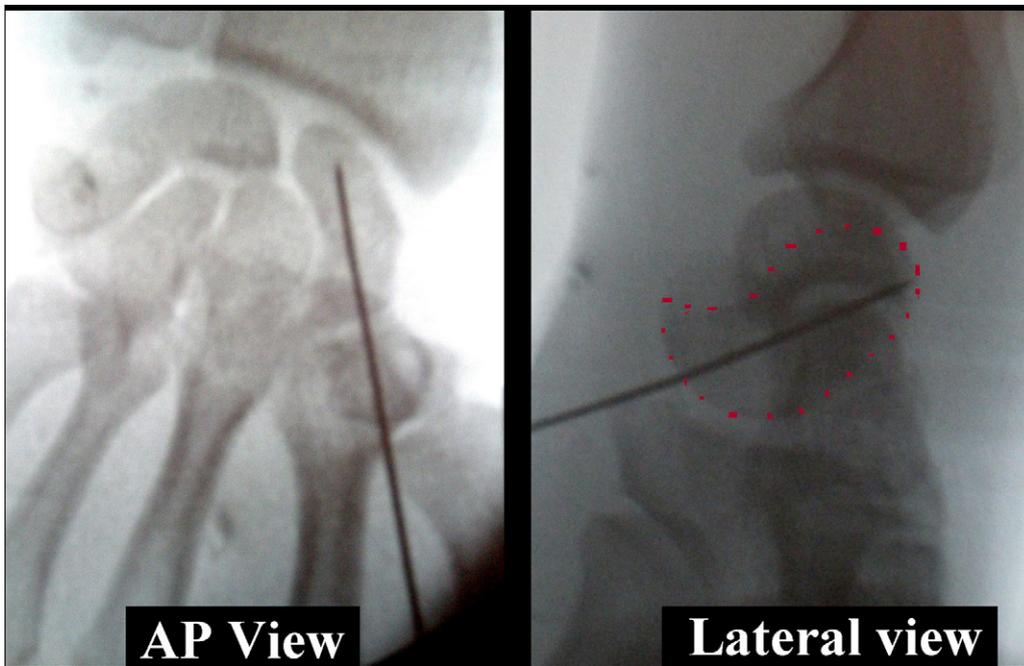


Fig. 2. An example of a guidewire correctly placed into the scaphoid.

104 *Technique with the GID (Figs. 3 and 4)*

105 The wrist was placed on the armboard under the image-
106 intensifier beam and locked in supination and extension using a

radiotransparent **wrist-stabilising** device (Chirobloc®, Arex, 107
Palaiseau, France). The **wrist-stabilising** device was attached to 108
the armboard with adhesive bands. The C-arm was free to rotate 109
from the vertical position to the horizontal position, which **was** 110

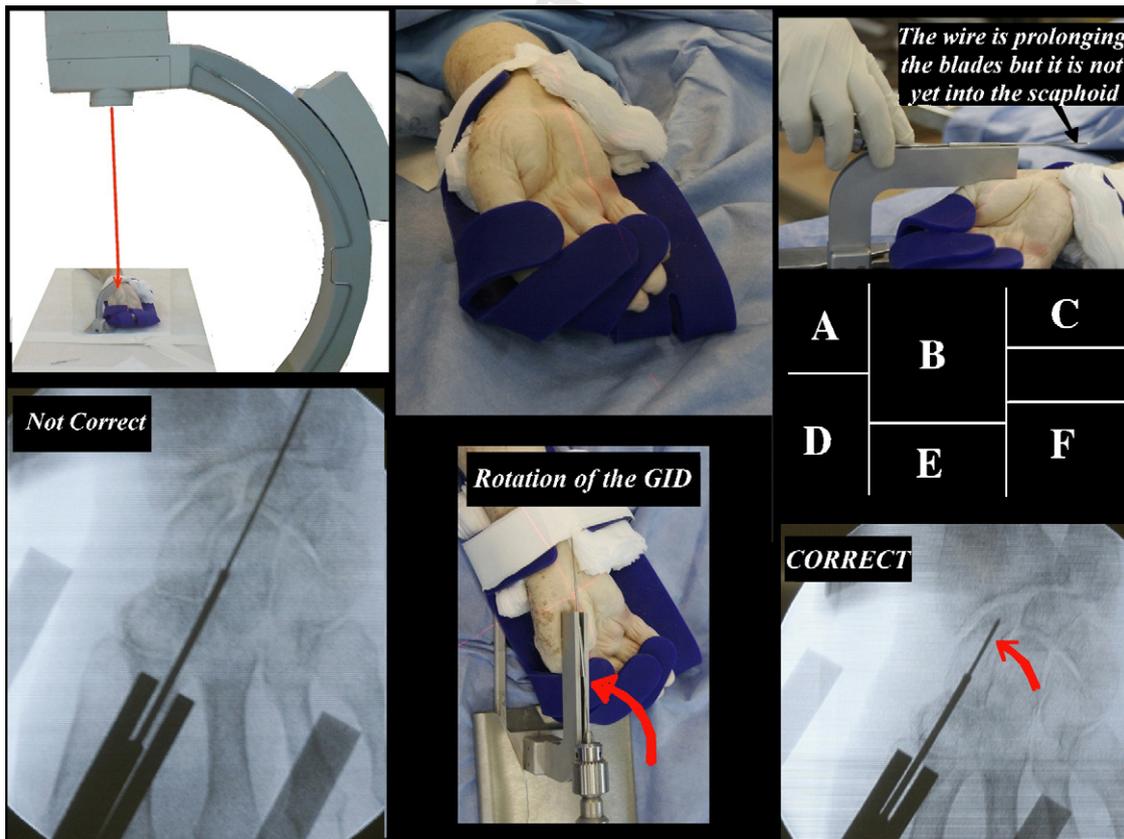


Fig. 3. The C-arm is vertical (A) and the wrist is **immobilised** with the radiotransparent handle (B). At this time, the wire is out of the wrist, prolonging the blades of the GID (C). The C-arm of the image intensifier is vertical in order to obtain an anteroposterior view of the scaphoid and radioscopic projection of the guidewire on the scaphoid (A and D). The ideal direction is obtained by rotating the GID horizontally on the armboard (E). When the direction is considered as correct on the anteroposterior view (F), the GID and the specimen's hand are left in the same position.

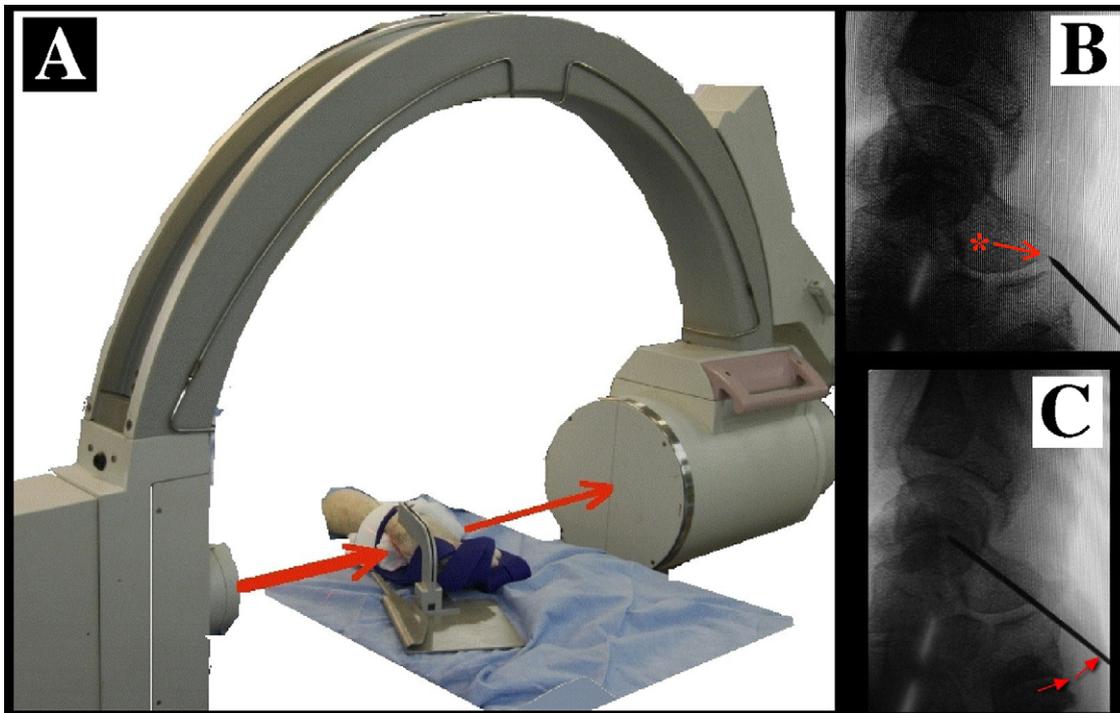


Fig. 4. The C-arm of the image intensifier is rotated horizontally (A) in order to obtain a lateral view of the scaphoid. The operator can move the guidewire vertically between the blades of the GID in order to find firstly the ideal entry point on the scaphoid distal pole (B) and secondly the optimal direction to reach the proximal pole (C). During all this step, the operator can move the wire between the blades of the GID without assessing the direction on the anteroposterior view which is locked by the GID. Thus, he can fully concentrate on the sagittal placement of the wire.

111 used to obtain anteroposterior and lateral views of the wrist,
 112 respectively. The procedure started with the C-arm of the image
 113 intensifier vertical to obtain anteroposterior views. The wire was
 114 introduced into the reducer, which was then inserted between the
 115 two blades of the guidewire device. The GID was moved
 116 horizontally until the guidewire projected over the scaphoid
 117 was in the desired position on the anteroposterior view. The **wrist-**
 118 **stabilising** device remained immobile on the armboard. Then, the
 119 C-arm was rotated 90° to obtain a lateral view without changing
 120 the position of the GID. On the lateral view, the wire was moved

121 vertically between the two blades of the GID until its tip projected
 122 on the base of the scaphoid tubercle. The wire was then aimed **at**
 123 **the centre** of the proximal pole and pushed into the scaphoid **with**
 124 **the drill**. Finally, the C-arm was rotated back to the anteroposterior
 125 view to assess wire position.

126 *Assessment of wire position in the scaphoid (Fig. 5)*

127 Each scaphoid was extracted by open dissection, leaving the
 128 wire into the bone. Computed tomography (CT) of each scaphoid

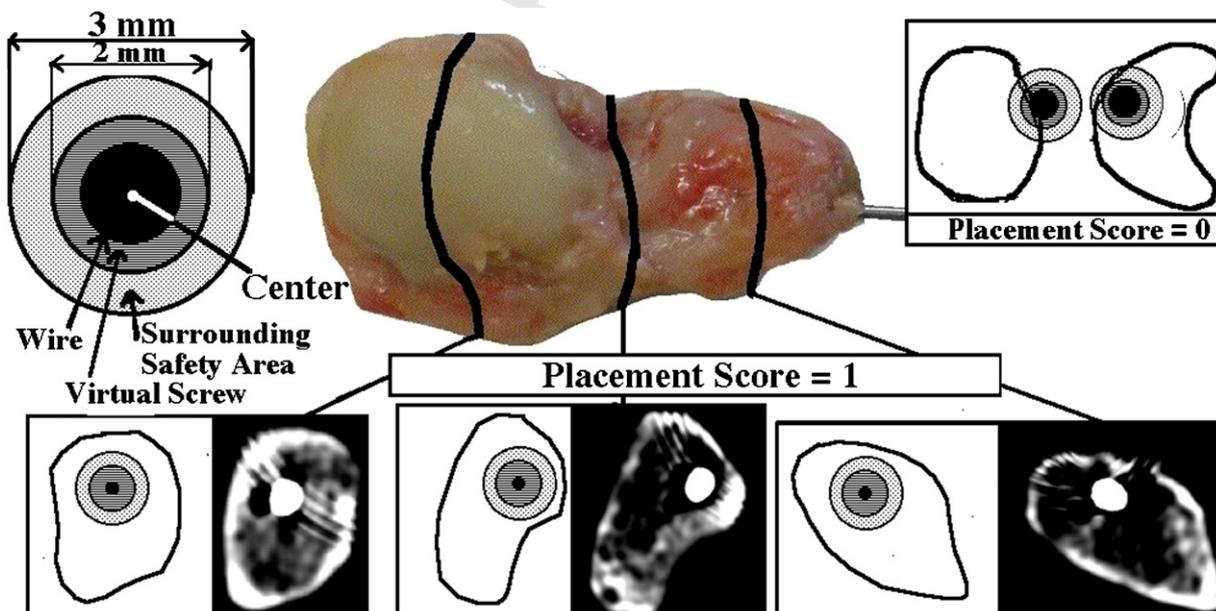


Fig. 5. Each scaphoid was extracted with the wire inside and was divided in three parts. A segmentary placement score (SPS) was calculated at each level. The global placement score (GPS) was the sum of the three SPS.

Table 1

Wrist	Specimen	Side	Surgeon	Technique	Time	Number of attempts	Number of image-intensifier shots	Distal SPS	Middle SPS	Proximal SPS	GPS
1	1	R	1	Hand held	723	8	31	1	0 (less than 2 mm with the scapho-capital joint)	1	2
2		L	1	GID	345	1	7	1	1	1	3
3	2	R	1	GID	306	1	9	1	1	1	3
4		L	1	Hand held	750	9	38	1	1	1	2
5	3	R	1	Hand held	624	5	22	1	1	0 (out of the radial surface of the scaphoid)	2
6		L	1	GID	282	1	6	1	1	1	3
7	4	R	1	GID	321	1	7	1	1	1	3
8		L	1	Hand held	587	6	30	1	1	1	3
9	5	R	2	Hand held	622	7	37	1	1	1	3
10		L	2	GID	362	1	5	1	0 (less than 2 mm with the scapho-capital joint)	1	2
11	6	R	2	GID	513	2	14	1	1	1	3
12		L	2	Hand held	382	4	22	1	1	1	3
13	7	R	2	Hand held	553	6	31	1	1	1	3
14		L	2	GID	334	1	5	1	1	1	3
15	8	R	2	GID	287	1	7	1	1	1	3
16		L	2	Hand held	375	4	18	1	0 (out of the scapho-capital joint)	1	2
17	9	R	3	Hand held	596	5	21	1	1	1	3
18		L	3	GID	320	2	12	1	1	1	3
19	10	R	3	GID	316	1	6	1	1	1	3
20		L	3	Hand held	503	5	23	1	1	0 (out of the radial surface of the scaphoid)	2
21	11	R	3	Hand held	356	4	22	1	1	1	3
22		L	3	GID	291	1	8	1	1	1	3
23	12	R	3	GID	298	1	7	1	1	1	3
24		L	3	Hand held	310	3	14	1	1	1	3
25	13	R	4	Hand held	367	4	21	1	1	1	3
26		L	4	GID	245	1	9	1	1	1	3
27	14	R	4	GID	270	1	7	1	1	1	3
28		L	4	Hand held	256	3	13	1	1	1	3
29	15	R	4	Hand held	301	2	11	1	1	1	3
30		L	4	GID	260	1	9	1	1	1	3
31	16	R	4	GID	275	1	7	1	1	1	3
32		L	4	Hand held	256	3	16	1	1	1	3

In 16 scaphoids, the guidewire was inserted by using the hand held technique and the GID was used for the 16 others. SPS: Segmentary placement score. GPS: Global placement score. GID: Guidewire insertion device.

was performed using a 16-slice multidetector scanner (Sensation 16®, Siemens Medical Solutions, Munich, Germany) to assess wire position. Slice reconstructions and measurements were done using Vitrea 2.2® software (Vital Image, Minneapolis, MN). For each scaphoid, three slices were reconstructed in planes perpendicular to the wire. Each slice was passing through the tubercle (distal third), the waist (middle third), and the middle of the radial surface (proximal third), respectively. To simulate the use of a 2-mm diameter screw, such as Herbert's screw, a circle 3 mm in diameter with the centre at the middle of the guidewire was drawn on each slice. Then, the segmentary placement score (SPS) was determined for each slice, as follows: the score was 0 if part of the circle was outside the scaphoid and 1 if the circle was entirely within the scaphoid. The global placement score (GPS) was calculated as the sum of the SPS values at the distal, middle, and proximal thirds.

Statistical analysis

Data are reported as medians with the interquartile range (IQR). The effect of using the guidewire device on procedure duration, number of placement attempts, and number of image-intensifier shots was assessed using multiple regression models accounting for surgeon experience. Multiple linear regression models accounting for heteroskedasticity and Poisson multiple regression models were built when appropriate. An interaction between surgeon experience and use of the guidewire device was looked for.

All analyses were performed using R 2.6.1.²⁷ All tests were two-sided, with the significance level set at 0.05.

Results

All the results are summarised in Table 1. Overall, median procedure duration was 338 s (IQR, 290-506), median number of attempts was 2 (IQR 1-3), median number of image-intensifier shots was 13 (IQR 7-22), and median GPS was 3 (IQR 3-3). With the hand held technique, median procedure duration was 443 s (IQR 345-603), median number of attempts was 4.5 (IQR 3.8-6), median number of image-intensifier shots was 22 (IQR 18-30), and median GPS was 3 (IQR 2-3). With the GID, median procedure duration was 314 s (IQR 280-324), median number of attempts was 1 (IQR 1-1), median number of image-intensifier shots was 7 (IQR 7-9), and median GPS was 3 (IQR 3-3).

The use of the GID significantly decreased procedure duration (P < 0.001), number of attempts (P < 0.001), and number of image-intensifier shots (P < 0.001). With both techniques, experienced surgeons were significantly faster (P = 0.0083) and required significantly fewer attempts (P = 0.043) than inexperienced surgeons. An interaction was found between the experience of the surgeons and the use of the GID. The GID had significantly greater effects in decreasing procedure duration (P = 0.0039) and number of image-intensifier shots (P < 0.001) for inexperienced surgeon than for the experienced surgeons. In contrast, no

177 interaction was found between surgeon experience and number of
178 attempts ($P = 0.32$).

179 Discussion

180 Acute nondisplaced fractures of the scaphoid waist are
181 common.^{18,20} PSF is an attractive option during which accurate
182 placement of the guidewire is often difficult to achieve. We
183 developed a GID to assist in guidewire placement. Compared to the
184 conventional **hand held** technique, the device significantly
185 decreased procedure duration, number of image-intensifier shots,
186 and number of placement attempts. Both experienced and
187 inexperienced surgeons obtained these benefits with the GID.

188 Correct wire placement is a prerequisite to successful PSF of the
189 scaphoid. The technical challenges raised by the placement of the
190 guidewire have prompted the development of guiding systems. In
191 their study describing the use of a double-threaded cannulated
192 screw in 158 patients, Herbert et al. recommended using a jig to
193 facilitate guidewire placement.¹⁶ However, the jig cannot be used
194 for percutaneous procedures. In a cadaver study of eleven
195 scaphoids, Liverneaux et al. found that fluoroscopic navigation
196 facilitated correct percutaneous guidewire placement, compared
197 to the conventional **hand held** technique.²¹ However, fluoroscopic
198 navigation is available only in **specialised centres**, whereas
199 nondisplaced scaphoid fractures are managed at all **centres**. The
200 GID we developed allows percutaneous procedures, and **does not**
201 need any other specific ancillary to be used. The radiopaque
202 **wrist-stabilising** device used in this study can be easily replaced by
203 an assistant keeping the hand in the desired position.

204 In our study, we compared the technique using the GID with the
205 "conventional" **hand held** technique. Meanwhile, there are two
206 different **hand held** techniques described in the literature and the
207 choice of the "conventional" technique of reference warrants few
208 justifications. The first **hand held** technique consists in leaving the
209 patient's hand freely movable under the beam of the image
210 intensifier.^{4,15} The second **hand held** technique uses chinese finger
211 traps to suspend the wrist which remains free to rotate under the
212 beam of the image intensifier.^{13,14} As Liverneaux et al.²¹ we **chose**
213 the first one as the **hand held** reference technique because it is the
214 one we usually perform in our department and because it is
215 probably the technique used by most of **the surgeons**.

216 Fluoroscopic guidance exposes the patient and surgeons to
217 **ionising** radiations, which can induce radiodermatitis and skin
218 cancer after several decades.^{22,30} Dose reduction during fluoro-
219 scopy-guided interventions is challenging.¹⁷ Some surgeons use
220 mini C-arm which allow minimal radiation exposure^{12,29} but that
221 kind of device remains restricted to a few number of **centres**
222 whereas most of orthopaedic surgeons in the world still use
223 conventional image intensifiers. For Shoaib et al., even if the mini
224 C-arm caused statistically less radiation to the surgeon, there was
225 no statistically significant difference in the radiation exposure to
226 the patient.²⁹ Moreover, even when using a mini C-arm, there is a
227 substantial amount of measurable radiation exposure for the
228 structures in line with the imaging beam.¹² Thus, radiation
229 exposure is unavoidable when performing PSF of the scaphoid
230 but the dose must be kept as small as possible. This was one of the
231 reasons why we developed the GID. A cadaver study showed that
232 fluoroscopic navigation was associated with a four-fold decrease in
233 radiation exposure, compared to the conventional wire-placement
234 method²¹ but nowadays, such a technology remains limited to a
235 few **centres**. Although the radiation dose varies with the type of
236 image intensifier, it is consistently proportional to the number of
237 shots. That is why we decided to use the number of **image-**
238 **intensifier** shots to estimate the amount of radiations. It would
239 have been tempting to use a dosimeter placed into the surgeon's
240 gloves and on the specimen's hand in order to measure the

delivered **dose more precisely**. Nevertheless, the measured data
would have been very dependant on the positioning (palmar or
dorsal side of the hands) and orientation of the dosimeters and it
would have been a potential source of confusion in data
interpretation. In our study, compared to the conventional **hand**
held technique, the use of the GID was associated with an about 3-
fold decrease in the number of **image-intensifier** shots, and
therefore in radiation exposure. In addition, the shorter procedure
duration and smaller number of attempts indicate facilitation of
wire placement with the GID.

Our study has several limitations. First, results obtained in
cadavers may not reflect in vivo results. However, the beneficial
effects of the device may be even larger in everyday practice,
when the restricted room available in the operating theater,
time constraints, stress, and fatigue may provide additional
opportunities for the device to facilitate wire placement. Second,
brief explanations about the device were given orally to the
surgeons before the study. Nonetheless, no attempts were
performed with the GID before the study (except by the first
author who developed the device) and all cases including the
first one were included in the analysis. Therefore, use of the GID
seems easy to learn. Second, our study does not demonstrate
that the GID is associated with improved clinical outcomes.
However, the decrease in radiation exposure is an important
benefit. Furthermore, easier wire placement may lead surgeons
to use PSF instead of prolonged cast **immobilisation** for treating
nondisplaced scaphoid fractures, which can be expected to
accelerate time to work resumption and time to recovery of
wrist function. For all these reasons, a clinical study already
started **assessing this GID in vivo**.

Conclusion

This cadaver study showed that the GID decreased the time and
radiation exposure needed to achieve correct percutaneous
guidewire placement in the scaphoid, compared to the conven-
tional **hand held** technique. These benefits were obtained both by
experienced and by inexperienced surgeons. Clinical studies of the
device are warranted.

Conflict of interest

None of the authors received fundings, grants, or in-kind in
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None of the authors have association or financial involvement
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